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A new species of death adder (*Acanthophis*: Serpentes: Elapidae) from north-western Australia

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Abstract

Australian death adders (genus *Acanthophis*) are highly venomous snakes with conservative morphology and sit-and-wait predatory habits, with only moderate taxonomic diversity that nevertheless remains incompletely understood. Analyses of mitochondrial and nuclear gene sequences and morphological characteristics of death adders in northern Australia reveal the existence of a new species from the Kimberley region of Western Australia and the Northern Territory, which we describe as *Acanthophis cryptamydros* sp. nov. Although populations from the Kimberley were previously considered conspecific with Northern Territory death adders of the *A. rugosus* complex, our mtDNA analysis indicates that its closest relatives are desert death adders, *A. pyrrhus*. We found that *A. cryptamydros* sp. nov. is distinct in both mtDNA and nDNA analysis, and possesses multiple morphological characteristics that allow it to be distinguished from all other *Acanthophis* species. This study further supports the Kimberley region as an area with high endemic biodiversity.

Key words: Australian Monsoonal Tropics, mtDNA, nDNA, systematics, taxonomy, Acanthophis cryptamydros sp. nov.

Introduction

The Kimberley region in north-western Australia is an area of high endemism (Slatyer *et al.* 2007; Bowman *et al.* 2010; Powney *et al.* 2010; Palmer *et al.* 2013; Pepper & Keogh 2014). Ongoing collections of animal and plant groups in recent years has led to the identification of many new species (e.g., frogs: Anstis *et al.* 2010; Doughty 2011; lizards: Doughty *et al.* 2012; Oliver *et al.* 2012, 2014; Pepper *et al.* 2011; snails: Köhler 2010, 2011; flowering plants: Barrett *et al.* 2009; Carlson *et al.* 2011; Maslin *et al.* 2013). Snakes have received some attention in the Kimberley region of Western Australia and the Australian Monsoonal Tropics (e.g. *Pseudechis*: Kuch *et al.* 2005; *Demansia*: Shea & Scanlon 2007; *Anilios*: Marin *et al.* 2013), yet many genera have not been subject to extensive systematic revision recently.

The death adders, *Acanthophis* Daudin, 1803 are a widespread genus of elapid snakes distributed across Australia and New Guinea, and on several Indonesian islands to the west of New Guinea. Although the genus is highly distinctive due to its convergent viper-like morphology and ecology (Shine 1980; Greer 1997), species limits within the genus have remained poorly understood (Storr 1981; McDowell 1984; Aplin & Donnellan 1999; Wüster *et al.* 2005), partly due to extensive polymorphism within many species and even populations (e.g., Johnston 1996).

The systematics of the death adders from the Australian Monsoonal Tropics (northern Queensland, the Top End of the Northern Territory and the Kimberley region in northern Western Australia) have proven especially complex. While *A. pyrrhus* Boulenger, 1898 from the arid zone is highly distinctive, and *A. wellsi* Hoser, 1998 from the Pilbara region in Western Australia was discovered, defined and shown to be a clearly diagnosable, valid

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species by Aplin & Donnellan (1999), the affinities of the remaining populations remain poorly understood. Storr (1981) assigned all populations from the Kimberley to *A. praelongus* Ramsay, 1877, an arrangement retained by many subsequent authors (e.g., Cogger 2000; Storr *et al.* 2002; Wilson & Swan 2013). Storr, however, indicated there was considerable heterogeneity in morphology within *A. praelongus senso lato*. More recent studies on venom also found considerable variation in composition across the range of *A. praelongus* (Fry *et al.* 2002). A molecular genetic study found that populations from the Northern Territory and north-western Queensland fell in to two distinct clades with partly overlapping distribution, with each more closely related to *A. rugosus* Loveridge, 1948 from New Guinea than to other Australian taxa (Wüster *et al.* 2005). These recent studies, however, did not include representatives from the Kimberley region of Western Australia. The Kimberley forms have been explicitly or implicitly grouped with Top End *Acanthophis* in the *A. rugosus* complex (they were previously considered part of *A. praelongus*). Storr (1981) did not distinguish Kimberley *Acanthophis* from Top End populations based on his morphological data and thus it has been assumed that they are conspecific with the *A. rugosus* complex (e.g., Cogger 2014). No specific comparison between these populations was made by Storr (1981); however, as at the time of his revision he only had access to 16 northern specimens in the collection of the Western Australian Museum, of which 13 were from the Kimberley region and three from the Northern Territory.

As part of a reanalysis of the systematics of *Acanthophis*, here we compare the Kimberley death adders to other *Acanthophis* using multiple molecular loci and morphology. We found the Kimberley specimens to be genetically and morphologically distinct from other currently-recognized taxa, and therefore describe this species as new below.

Material and methods

Molecular phylogenetics. Appendix 1 shows the specimen data for individuals and samples that were used for molecular phylogenetic analysis. Genomic DNA (ventral scale clippings, liver, and blood samples) was extracted from *Acanthophis* specimens using a Qiagen DNeasyTM Tissue Kit. NADH dehydrogenase subunit 4 (nd4) and cytochrome b (cytb) sequences from the Wüster et al. (2005) study were obtained from GenBank. Sampling localities are shown in Fig. 1. Based on the relationships presented by Sanders et al. (2008), two species were used as outgroups: *Pseudechis papuanus* (the likely sister genus to *Acanthophis*) and *Oxyuranus scutellatus* (a more distant outgroup).

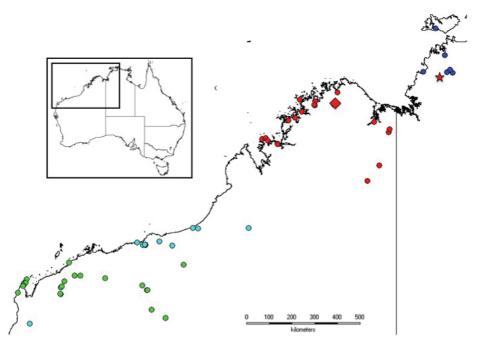


FIGURE 1. Distribution of *Acanthophis* sampled in northwest Australia. Only samples with accurate collection coordinates have been included, except specimen NTM R29109 (star; see text). Colored circles correspond to sampled specimens: red = A. *cryptamydros* **sp. nov.**; blue = A. *rugosus*; turquoise = A. *pyrrhus*; green = A. *wellsi*. The collection locality of the holotype of A. *cryptamydros* (WAM R174083) is displayed as a red diamond.

Three mitochondrial gene fragments (nd4, cytb, and 16s) and two nuclear gene fragments (prolactin receptor (prlr) and ubinuclein 1 (ubn1)) were amplified using the polymerase chain reaction (PCR). Primer information is shown in Table 1. PCR reaction volume was 11 μ l, which consisted of 9.6 μ l of Abgene 1.1x ReddyMixTM (1.25 units Thermoprime Plus DNA polymerase; 75 mM Tris-HCl pH 8.8; 20mM (NH₄)₂SO₄; 1.5 mM MgCl₂; 0.01% (v/v) Tween®20; 0.2 mM of each dNTP; and a precipitant red dye for electrophoresis), 0.3 μ l of required primer and ~10 ng/ μ l of template DNA per sample. Amplification conditions for reactions was denaturation at 94°C for 2 minutes (min); then 35 (16s, ubn1 and prlr) or 40 (nd4 and cytb) cycles of 94°C for 30 seconds (s); annealing at 43°C (16s), 48°C (16s), 50°C (16s),

Sequences were proofread and aligned using CodonCode Aligner 3.7.1 using default settings and checked for pseudogenes, unexpected stop-codons or indels using Molecular Evolutionary Genetics Analysis 4.0.2 (MEGA) (Tamura *et al.* 2007) by translating the DNA sequences into amino acid sequences.

For phylogenetic analysis, Bayesian inference (BI) methods were applied to the mtDNA dataset and haplotype networks were applied to the two nuclear genes separately to infer phylogenetic relationships. BI analysis was implemented through MrBayes 3.1.2 (Ronquist & Huelsenbeck 2003). Coding genes were partitioned and the most suitable model of sequence evolution under the Akaike Information Criteria (AIC) for each gene and partition was identified using MrModeltest 2.3 (Nylander 2004), implemented in PAUP* 4.0b10 (Swofford 2002). MrBayes analyses ran for 10⁷ generations, with two independent parallel runs with one cold and three heated chains each. Trees were sampled every 1000 generations and the first 10% of trees were discarded as burn-in. Effective sample size and burn-in for each parallel run were examined in Tracer v.1.5 (Rambaut & Drummond 2007). Analysis was run through the Bioportal (http://www.bioportal.uio.no/). Maximum likelihood (ML) analysis was also carried out on the partitioned mtDNA dataset using the program Randomized Axelerated Maximum Likelihood (RAxML) 7.2.2 (Pfeiffer & Stamatakis 2010) with 500 bootstrap replicates.

TABLE 1. Primers used during PCR. ^aArévalo *et al.* (1994); ^bWüster *et al.*(2008); ^cnewly developed primers; ^dTownsend *et al.* (2008); ^cAxel Barlow (pers. comm.).

Primer Name	Primer $(5'-3')$	
nd4		
ND4 ^a	CACCTATGACTACCAAAAGCTCATGTAGAAGC	
H12763V ^b	TTCTATCACTTGGATTTGCACCA	
cytb		
AcanF2°	CTACTATCCTCCAACCT	
AcanR1°	CAGTTTGTTTGGGATTGATCG	
16s		
16sL ^a	CGCCTGTTTATCAAAAACAT	
16sR ^a	CCGGTCTGAACTCAGATCACGT	
prlr		
PRLR_F1 ^d	GACARYGARGACCAGCAACTRATGCC	
PRLR_R3d	GACYTTGTGRACTTCYACRTAATCCAT	
ubn1		
BaUBN_F°	CCTCTGGTTACTCAGCAGCA	
BaUBN_R°	ATTGGCCACTCCTTGTGTTC	

seqPHASE (Flot 2010) was used prior to the implementation of PHASE 2.1.1 (Stephens *et al.* 2001; Stephens & Scheet 2005) to infer nuclear haplotypes when heterozygous positions were present in the dataset. Heterozygous positions were considered to be true if they received a probability of >0.7. Resulting sequences were checked for indels or stop codons in MEGA prior to producing haplotype networks in Network 4.6.0.0 using maximum parsimony criteria.

In order to assess patterns of genetic variation across all three loci together (the two mtDNA genes were combined and treated as a single locus), we used the program POFAD 1.03 (Joly & Bruneau 2006) to generate standardized multilocus distances between specimens. POFAD allows the incorporation of allelic variation, and is thus very useful when assessing genetic distances between specimens in nuclear loci. Moreover, it allows equal weighting of all loci, rather than allowing the more variable loci to dominate the analysis. Only individuals that had all loci successfully amplified were used in the analysis and MEGA was used to produce the data matrices of *p*-distances for each gene, which were generated using pairwise deletion of missing data. Due to its nature, the *ubn1* dataset contains indels spanning two separate codons. MEGA treats such data as missing and thus the *p*-distance would be 0. To overcome this problem, a single nucleobase for each indel was inserted into the end of each sequence with those containing the indel having a different nucleobase coded for them compared to those without. The resulting standardized between-specimen multilocus distance matrix was visualised with a principal coordinates analysis (PCO) using the MVSP program (www.kovcomp.com).

Morphological analysis. Appendix 1 shows specimen data for all specimens, including type material, examined for morphological comparisons from the collections of the Western Australian Museum (WAM) and Museum and Art Gallery of the Northern Territory (NTM). We used 14 genotyped specimens from the phylogenetic analysis and a further 86 non-genotyped specimens to identify consistent morphological characteristics between the new species and other Australian *Acanthophis* species, including those within the *A. rugosus* group.

Table 2 presents the morphological variables measured. Characters examined for the morphological analysis included body measurements, scale counts, and the size, shape, and positioning of head scales of each specimen. Measurements, scale counts, and morphological nomenclature follow that of Storr *et al.* (2002) and Cogger (2014). There were missing scores for some specimens due to poor condition, such as road mortalities and very old or poorly preserved specimens.

TABLE 2. Meristic and morphological characters measured in this study.

Character	Description
ToL	Total body length, from snout to tail tip, excluding terminal spine
SVL	Snout-vent length, from anteriormost point of snout to posterior edge of anal scale
TailL	Tail length, from anterior edge of first subcaudal scale to tail tip, excluding terminal spine
HeadL	Head length, from tip of snout to posterior margin of the of the quadrate
HeadW	Head width, at widest point, posterior to eyes
VS	Ventral scales, counted from anteriormost ventral to but not including anal scale
MBSR	Midbody scale rows, at middle of SVL
AntSR	Anterior scale rows, posterior to head
PostSR	Posterior scale rows, anterior to vent
ScST	Subcaudal scales total, from the first subcaudal posterior to vent to posterior most scale on tail tip, including single and paired scales, excluding terminal spine
ScSS	Subcaudal scales single, as for ScST, excluding paired ScS
ScSP	Subcaudal scales paired, as for ScST, excluding single ScS
FrL	Frontal length, from anteriormost edge to posterior most edge
FrW	Frontal width, at widest point
SupOcL	Supraocular length, from anteriormost edge to posterior most edge
SupOcW	Supraocular width, at widest point
SupLab	Supralabial scales
InfLab	Infra labial scales

SVL and TailL were measured with a ruler to the nearest 1.0 mm. TailL is presented as length (mm) and as a percentage of SVL. All other meristic variables including HeadL and HeadW, FrL and FrW, and SupOcL and SupOcW were measured with digital calipers to the nearest 0.1 mm. Head and selected scale meristics are presented as length and width (mm) and width as a percentage of length. MBSR counts were taken at approximately 50% of the SVL. Due to variation in the number of scale rows along the length of the body, often reducing anteriorly and posteriorly, multiple counts were taken to determine the minimum number of scale rows. Scale row counts were also taken from posterior to the head (AntSR), and anterior to the vent (PostSR). VS were counted from the anteriormost ventral to, but not including, the anal scale following the Dowling (1951) method. Subcaudal scale counts were taken from the first subcaudal posterior to the vent to the terminal scale on the tail tip, excluding spine. Where possible, the sex of specimens was determined from everted hemipenes, presence of follicles or eggs, or by direct examination of reproductive organs via ventral incision.

Color in life was based on recently collected specimens and photographs of specimens in life. In addition to the dorsal color and pattern, the coloration and extent of pigmentation on the ventral surface of specimens was also examined.

Results

Molecular genetics. A total of 2,874 bp were used in phylogenetic analysis: 684 for *nd4* (169 were variable and 153 were parsimony informative); 752 for *cytb* (179 were variable and 161 were parsimony informative); 473 for *16s* (53 were variable and 26 were parsimony informative); 508 for *prlr* (19 were variable and 13 were parsimony informative); and 457 for *ubn1* (19 were variable and 9 were parsimony informative).

The consensus Bayesian inference (BI) tree is shown in Fig. 2 with maximum likelihood (ML) scores mapped on. The Kimberley *Acanthophis* clade is closest genetically to *A. pyrrhus* in the mitochondrial phylogeny, with a mean *p*-distance of 7.4%. The rest of the phylogenetic relationships are consistent with those presented by Wüster *et al.* (2005), and are summarized as follows. Sister to these two lineages is the Pilbara endemic *A. wellsi*. These three lineages form a well-supported sister group to the remaining Australian populations (i.e., *A. antarcticus*, *A. praelongus*, and an '*A. rugosus* group' comprised of multiple more weakly-diverged lineages). *Acanthophis laevis* from New Guinea forms the sister taxon to all other species.

The nDNA data further support the Kimberley population as an independently evolving lineage (Figs. 3 and 4), with only a small amount of haplotype sharing occurring in both *prlr* and *ubn1* (Fig. 3). In *prlr* there are two haplotypes shared between *A. rugosus* and Kimberley specimens. In both *prlr* and *ubn1* one haplotype is shared between *A. wellsi* and specimens from the Kimberley. Specimens from Kimberley, however, also contain multiple unique haplotypes; six in *prlr* and five in *ubn1*. The principal coordinate analysis of standardized multilocus distances supports the distinction of the Kimberley population compared to other closely related species (Fig. 4).

Morphology. Table 3 summarizes the meristic differences among the forms examined. In terms of overall body length and proportions, the Kimberley specimens were relatively homogeneous, and broadly overlapped with specimens from the *A. rugosus* group and other Australian *Acanthophis* species. More apparent, however, were differences in quantitative characters, scalation, and pattern that corresponded with lineages identified from the molecular analyses revealing diagnosable groups. Populations of *Acanthophis* from the Kimberley region of Western Australia and western Northern Territory differ from lineages with the *A. rugosus* group by possessing the following characteristics (see below for comparisons with other species): 22 or 23 MBSR; 125–139 VS; undivided prefrontal scales; posterior edge of frontal scale not extending beyond posterior edge of supraoculars; laterally flared supraoculars; area of lower secondary temporal scale equal to or smaller than sixth supralabial; anterior dorsal scales with prominent keels and unpigmented ventrum except for 1–3 rows of spots on ventrolateral edge.

Systematics

Our mtDNA results generally conform to the topology presented by Wüster *et al.* (2005). Contrary to the traditional grouping of the Kimberley populations with the '*Acanthophis rugosus*' group, however, the Kimberley population (Fig. 2) is the sister of *A. pyrrhus*, with *A. wellsi* as the sister taxon to this group. Moreover, each taxon

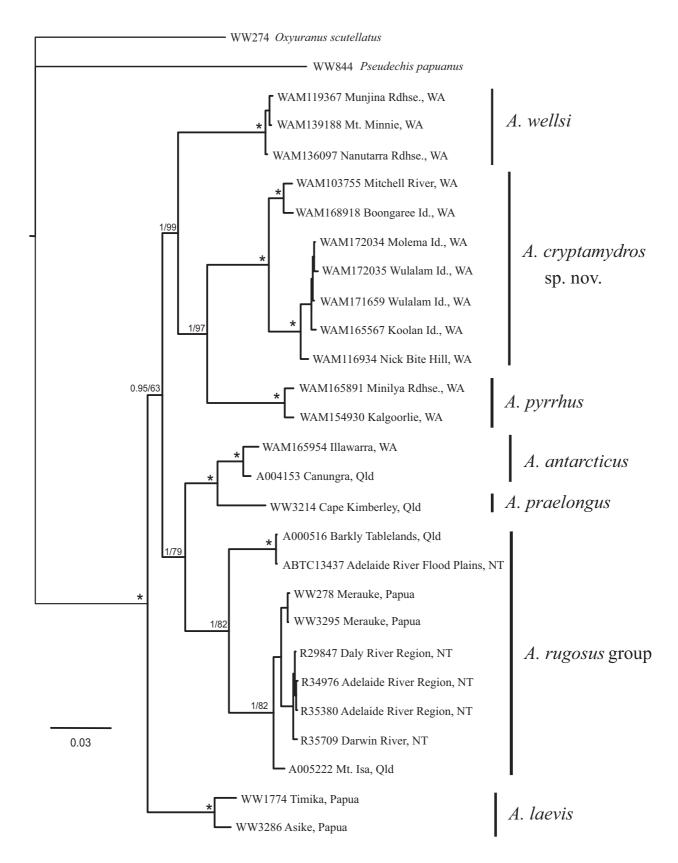


FIGURE 2. Mitochondrial phylogeny of *Acanthophis* inferred from Bayesian inference. Numbers at nodes indicate Bayesian posterior probability and maximum likelihood bootstrap supports, respectively. *s on nodes indicate maximum support for BI and ML.

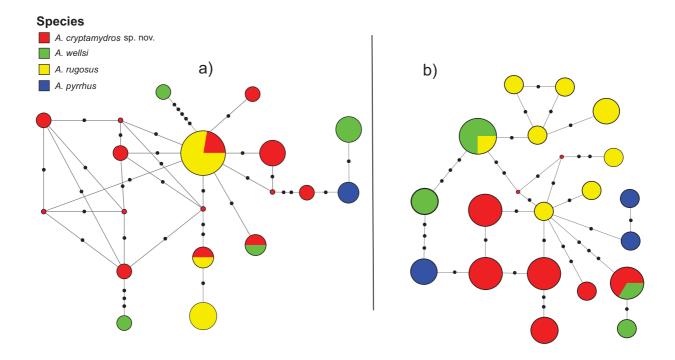


FIGURE 3. Haplotype networks showing relationships between *A. cryptamydros* **sp. nov.**, *A. wellsi*, *A. rugosus*, and *A. pyrrhus* for the two nuclear genes: a) *prlr*; and b) *ubn1*. Black circles indicate median vectors. Size of haplotypes is proportional to the number of specimens containing that haplotype.

(except A. pyrrhus and A. wellsi in ubn1) is distinguished by possessing one or several unique haplotypes not shared with other taxa. In particular, the Kimberley specimens show distinctive haplotype assemblages (Fig. 3), with multiple unique haplotypes not shared with other species, in both nuclear genes sampled. This is also reflected in patterns of overall genetic differentiation across all three loci, in which the Kimberley population forms a distinct but cohesive cluster (Fig. 4).

Morphological analysis found several characteristics that consistently diagnose the Kimberley population from all other Australian *Acanthophis* species (see above). Although characters in *Acanthophis* can vary widely and do show overlap with values in other species, the Kimberley taxon is reliably diagnosed in most cases by using a combination of scale characteristics and color pattern.

The consistent differences between the Kimberley death adders and all other *Acanthophis* across three independent genetic loci, morphology, and color pattern lead us to conclude that these populations represent a separate species from all other Australian *Acanthophis*. Since the only existing name applicable to this taxon, *Acanthophis lancasteri* Wells & Wellington, 1985, is a *nomen nudum* (Aplin & Donnellan 1999), we describe it as a new species below, diagnosing it from its congeners and all other currently recognized Australian *Acanthophis* species.

Elapidae

Acanthophis Daudin, 1803

Type species. Boa antartica (= Acanthophis antarcticus) Shaw & Nodder, 1802, by monotypy.

Diagnosis. Species assigned to the genus *Acanthophis* are moderately large, stout terrestrial elapid species most similar to vipers (Viperidae) from other continents. Species within the genus have distinctive wide and stout heads anterior to a defined narrowing forebody that rapidly broadens to the widest point towards midbody. Tail slender, distal portion laterally compressed terminating in a tail spine.

Etymology. From the Greek words *acanthi* meaning 'spine' and *ophis* meaning 'snake', in reference to the terminal tail spine present on species within the genus.

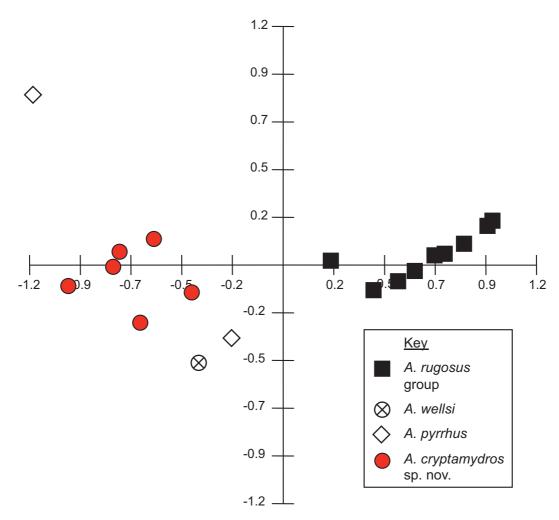


FIGURE 4. Principal coordinates analysis of *Acanthophis* taxa considered here, based on mean *p*-distance of all three loci used herein, with equal weighting.

Acanthophis cryptamydros sp. nov.

Kimberley death adders

Figs. 5–8

Holotype. WAM R174083, medium-sized male collected 1 km north-west of Theda Station homestead, Western Australia (14°46'59.10"S, 126°29'22.02"E), on 8 March 2014 by R. Ellis, G. Bourke, and R. Barrett. Fixed in 10% formalin, stored in 70% ethanol at WAM. Liver samples stored in 100% ethanol at WAM and SAM.

Paratypes. WAM R70690, sub-adult male, 45 km north-northeast Halls Creek, WA (17°51'00"S, 127°50'00"E); WAM R81245, adult male, Packsaddle Springs, near Kununurra, WA (15°54'00"S, 128°41'00"E); WAM R103755, adult male, Surveyors Pool, Mitchell River National Park, WA (14°39'46"S, 125°44'34"E); WAM R165567, adult male, Koolan Island, WA (16°08'04"S, 123°45'05"E); WAM R168918, adult female, Boongaree Island, WA (15°4'39.36"S, 125°11'13.56"E); WAM R172034, adult female, north-west Molema Island, WA (16°14'17"S, 123°49'49"E).

Diagnosis. A moderately stout *Acanthophis* to 645 mm total length. Distinguished from all other Australian *Acanthophis* by a combination of midbody scales in 22 or 23 rows, 125–139 ventrals, undivided prefrontal scales, posterior edge of frontal scale not extending beyond posterior edge of supraoculars, laterally flared supraoculars, area of lower secondary temporal scale equal to or smaller than sixth supralabial, anterior dorsal scales with prominent keels, and ventrum unpigmented except for 1–3 rows of spots on ventrolateral edge.



FIGURE 5. Acanthophis cryptamydros sp. nov. holotype (WAM R174083) in life (photograph—R.J. Ellis).

Description of holotype (WAM R174083). A medium-sized male *Acanthophis*, measurements and counts: ToL 482 mm; SVL 394 mm; TailL 88 mm (22% of SVL); HeadL 24.9 mm; HeadW 17.4 mm; MBSR 23; AntSR 20; PostSR 17; VS 129; ScST 54; SupLab 6; InfLab 7.

From above, head pear-shaped and distinct from neck, widest at interparietal scale angling forward to rostral and back to posterior jaw edge, narrowing to body; tip of snout blunt, broadly rounded; head in profile deepest at interparietal scale, snout convex; top of snout slightly concave where rostral and internasals converge; head scales rugose; rostral scale twice as wide as high, apex rounded, distal edges with low straight sides, ventral edge concave above lingual fossa; nasal scales narrowly separated by two internasals, approximately twice as wide as tall, rugose; nostril centered on nasal scale, opening dorsally and posteriorly, a shallow divot posterior to nostril; internasals as wide as long, in broad contact with each other, narrow contact with rostral, broad contact with nasals and prefrontals; prefrontals 1.5 times long as wide, narrowing laterally, 0.75 times area of frontal, 1.5 times area of internasals; frontal scale roughly rectangular, anterior edge slightly wider than posterior edge, approximately 1.5 times as long as wide (FrL 4.7 mm, FrW 3.0 mm), apex of posterior edge not extending beyond posterior edge of supraoculars; two parietals, as wide as long, anterior edge in contact with posterior angles of frontal, anterior edge of parietal sharing border with posterior edge of frontal, anterolateral edge in broad contact with supraocular, narrow contact with upper postocular, posterior border irregularly scalloped; preocular single, supraocular single, postoculars two, suboculars two; preocular in contact with prefrontal, nasal, third supralabial and anterior subocular scales; supraocular much longer than wide (SupOcL 5.03 mm, SupOcW 2.90 mm), thicker and rugose in appearance to other head scales, angled upwards at 30°; primary temporal scales two, lower primary temporal 2.5 times larger than upper, upper primary temporal feebly keeled; secondary temporal scales four, gradually increasing in size from dorsalmost to ventralmost, first and second with moderate keels, third and fourth smooth, fourth secondary temporal larger than prior three, located in the notch formed by fifth and sixth supralabials, two times smaller than sixth supralabial; supralabials six, sixth largest, fifth slightly smaller; first supralabial in contact with rostral and nasal, second in contact with nasal; third in contact with nasal, preocular, primary subocular, secondary subocular; fourth in contact with secondary subocular and lower postocular; fifth in contact with lower

subocular, second primary temporal, fourth secondary temporal and sixth supralabial; mental triangular; infralabials seven, fourth infralabial largest, first in contact with postmental scale; anterior chin shields in contact with infralabial only, anterior and posterior chin shields forming a butterfly-like shape; six rows of intergulars between chin shields and anteriormost ventral.

Body width widest at midbody tapering gradually forward to base of head and posteriorly to cloaca; scale rows 23 at midbody (i.e., at 64th ventral scale from anterior), decreasing to 20 behind the head, 17 anterior to vent; 129 ventral scales; anal scale single; 54 subcaudal scales, first subcaudal paired, anteriormost 29 single, followed by 29 paired; scales on side of body diamond-shaped, scales in vertebral zone more narrow; dorsal keeling strongest on anterior quarter of body, 10–12 longitudinal scale rows wide; keeling weak along remainder of length; dorsolateral and lateral scales weakly keeled to smooth, ventral scales smooth (Figs. 5–7).

Tail elongate, TailL 88 mm (22% of SVL), tapering from cloaca to laterally compressed caudal lure; ScST 54 (ScSS 30, ScSP 24). Caudal lure much higher than wide ending with terminal tail spine.

Eyes small with vertically elliptic pupil, iris mottled in appearance, similar in coloration to surrounding ocular scales.

Coloration. In life, ground color of dorsal and lateral surfaces pale orange-brown; 33 darker cross-bands (Fig. 5); cross-bands two to four midbody scales wide with dark brown border; dorsal scales edged with black anteriorly; tail coloration same as dorsum with 12 bands; tail tip and terminal spine black with white to cream flecks; first midbody scale dark in center, distal edge pale; ventral scales cream-white and lacking pigmentation other than lateralmost edges; ventral coloration of tail similar in appearance to ventrum with patches of dark pigment in center of posteriormost subcaudal scales prior to black tail tip and terminal spine; supralabials stippled with black, stronger stippling on posterior labials, ventral edge pale white; infralabials pale white with dark oblong vertical blotch in center of scale, on posterior scales the ventral edge of the blotch is angled posteriorly.

In preservative, dorsal ground color dull orange brown; overall pattern is subdued with less contrast between light and dark cross-bands (Fig. 6).



FIGURE 6. Acanthophis cryptamydros sp. nov. holotype (WAM R174083) showing dorsal and ventral coloration and pattern.



FIGURE 7. Head scalation and patterning of Acanthophis cryptamydros sp. nov. holotype (WAM R174083).



FIGURE 8. Variation in *Acanthophis cryptamydros* **sp. nov.** WAM R174083 (holotype), NTM R29109 and WAM R172034 (paratype).

Variation. SVL up to 555 mm; TailL 15–24% of SVL, mean 20% (N = 22). FrW 47–73% of FrL, mean 59% (N = 26). Apex of posterior edge terminates equal to (N = 13) or prior to (N = 13) posterior supraocular edge, never post. Fourth secondary temporal equal to (N = 13) or smaller than (N = 13) sixth supralabial. MBSR 22–23 (N = 24), mostly 23 (N = 21), occasionally 22 (N = 3). AntSR 16–23, mean 19 (N = 24), PostSR 16–19, mean 18 (N = 24). Ventral scales 125–139, mean 130 (N = 23). First anterior subcaudal scale usually divided, not separated (N = 9), occasionally undivided (N = 5), divided and separated by two (N = 4), three (N = 3) or one (N = 1) by a small rounded scale. ScST 46–56, mean 50, ScSP 20–40, mean 30 and ScSS 14–32, mean 21 (N = 22).

Ground color of dorsal and lateral surfaces variable from dull red-orange, tan-brown or gray in coloration with darker cross bands (Fig 6). Dorsal cross bands 44–61, mean 50 (N = 23); SVL 35–46, mean 39, Tail 9–15, mean 11 (N = 23). Tail tip black with white ventral surface, occasional small white lateral flecks (80%, N = 16), less often white (15%, N = 3) or banded (5%, N = 1). Ventral scales cream-white and lacking pigmentation other than lateral-most edges of ventral scale. Supralabials pale white, mottled in appearance fusing to darker markings on adjacent scales, lower edge of supralabials pale. Infralabials white-edged with dark pigment, both solid and mottled in appearance in center of scale extending to upper edge of scale.

Sexual dimorphism is not obvious, although female TailL % of SVL is shorter than males: female TailL 15–19% of SVL, mean 18 (N = 10); male TailL 21–24% SVL, mean 22% (N = 10). The tail tapers much more abruptly posterior to the vent in females, whereas in males the tail tapers gradually. The number of ventral and subcaudal scales is similar in both sexes (Table 3).

Distribution. Acanthophis cryptamydros **sp. nov.** is known from the Kimberley region of Western Australia. The species' range in Western Australia is known to extend from Wotjulum (WAM R11241) in the west, 45 km north-north-east of Halls Creek in the south (WAM R70690), and Kununurra in the east (WAM R137470). Acanthophis cryptamydros **sp. nov.** is also known to occur on some offshore islands including Koolan, Bigge, Boongaree, Wulalam, and an unnamed island in Talbot Bay (Palmer *et al.* 2013).

A single specimen (NTM R29109) with incomplete data (no collection date or precise latitude and longitude) is reported as occurring from Adelaide River in the Northern Territory (Fig. 1); however, this locality may be in error. Further collecting from the area may resolve the issue.

Habitat and ecology. The holotype (WAM R174083) was collected early morning towards the end of the 'wet season' (early March) from among basalt boulders in savannah woodland at the edge of a low plateau near a small sandstone outcrop. Vegetation among the basalt boulder habitat was dominated by *Eucalyptus tectifica* and *Corymbia greeniana* amongst mixed shrubs over mixed groundcovers and annual/perennial grasses (Fig. 9). Shrub cover was dominated by *Grevillea mimosoides*, *Grewia retusifolia*, *Indigofera* sp. and *Olearia arguta* (Fig. 9). The specimen was observed retreating to grass tussocks after being disturbed. The specimen was tightly coiled within the grass tussock before attempting to move to another tussock when disturbed. WAM R145216 was collected from thick grasses on a creek bank subjected to minor sheet flooding at Little Mertens Falls. Collection notes and accession data of other specimens describe individuals collected from among grasses or leaf litter in sandstone habitats. One specimen was collected from under a rock in a vine thicket near a beach on an unnamed island in Talbot Bay (WAM R172034), and another from leaf litter in *Acacia* woodland on Wulalam Island (WAM R172035). Two specimens were collected from roads or tracks (WAM R70690, WAM R165567).

Examination of the stomach contents of two specimens revealed a mixture of frogs, lizards, and mammals. A juvenile specimen (WAM R1251B) had an adult *Heteronotia* species (aff. *H. binoei*). In the gut of an adult specimen (WAM R141552), a subadult *Lophognathus* species (*L. gilberti*), an adult *Litoria* species (aff. *L. nasuta* or *L. watjulumensis*), and hair belonging to a native murid species, *Pseudomys* sp. (aff. *P. johnstoni* or *P. delicatulus*) were found. Two small reptilian eggs were also collected from the adult, possibly from the *Lophognathus* ingested. Accession data for WAM R116934 identified a *Ctenotus pantherinus* (WAM R117001) from examination of its stomach contents. An ecological study of Australian and Papuan death adders by Shine *et al.* (2014) included five specimens of *A. cryptamydros*; however, examination of stomach contents revealed no prey items, only a small quantity of dirt. Examination of the gut contents of other northern *Acanthophis* species indicated a diet consisting of a wide range of vertebrate species, especially lizards but also including frogs, mammals, and some birds (Shine *et al.* 2014).

Examination of reproductive organs of an adult female specimen (WAM R106033) revealed 13 well-developed follicles approximately 10 x 14 mm in size (month of collection unknown).

Comparison with other species. Distinguished from *A. pyrrhus* by higher average MBSR (23 vs. 21), fewer ventral scales (125–139 vs. 136–158), pigment on lateral periphery of ventral scales (vs. no pigment on ventral scales), presence of pigment patches on infralabials (infralabials unpigmented in *A. pyrrhus*), less prominent dorsal keeling (strongly keeled in *A. pyrrhus*, tapering to a sharp point on posterior edge), head scales less rugose, posterior edge of frontal scale not extending beyond posterior edge of supraoculars (equal to or beyond in *A. pyrrhus*), and undivided pair of prefrontal scales (divided in *A. pyrrhus*).

TABLE 3. Summaries of characters and ratios measured for *A. cryptamydros* **sp. nov.** and *A. 'rugosus* group'. All measurements in mm. Mean±S.D. (range). See Table 2 for abbreviations. Sample size listed in column headings, unless noted for individual characters below. Damaged specimens where accurate counts or measurements could be obtained or those less than 300 mm SVL were excluded from measurements and ratios but included in scale counts.

	A. cryptamyo	dros sp. nov.	A. 'rugos	us group'
Character	N =	: 23	N =	= 14
	(9♂,	7♀)	(9♂,	5♀)
SVL	428±57 (322-	-555) N = 20	486±97 (375–690)
	$\sqrt[3]{N} = 9$	$\mathcal{P} \mathcal{P} N = 7$	$\sqrt[3]{N} = 9$	Q Q N = 5
	412±41	468±54	434±57	581±82
	(374–496)	(392–555)	(375–530)	(482–690)
TailL	86±9 (72–1	10) $N = 20$	94±11 (80–110)
	$\sqrt[3]{3}$ $N = 9$	$\mathcal{P} \mathcal{P} N = 7$	$\sqrt[3]{N} = 9$	
	90.6±9	81.4±5	92±11	98±9
	(79–110)	(76–89)	(79–110)	(85–108)
TailL/SVL	0.20±0.03 (0.1	5-0.24) N = 20	0.20 ± 0.03	(0.15–0.23)
	$\sqrt[3]{N} = 9$		$\sqrt[3]{N} = 9$	
	0.22 ± 0.01	0.18 ± 0.01	0.21 ± 0.01	0.17 ± 0.03
	(0.21-0.24)	(0.15–0.19)	(0.20-0.23)	(0.15-0.22)
HeadL	26.1±2.4 (22.7	(-32.0) N = 20	31.1±5.5 (25.5–42.3)
HeadW	16.4±2.2 (13.8	(-20.2) N = 20	19.4±4.6 (13.1–27.1)
HeadW/HeadL	0.63±0.05 (0.54	4-0.71) N = 20	0.60 ± 0.04	(0.51–0.66)
VS	130±4 (125-	-139) N = 20	127±4 (1	23–136)
ScST	50±3 (46-	56) $N = 20$	46±6.14	(29–55)
	$\sqrt[3]{3}$ N = 9	$\mathcal{P} \mathcal{P} N = 7$	$\sqrt[3]{N} = 9$	
	53±2	49±2	47±7	45±3
	(51–56)	(46–51)	(29–55)	(42–49)
MBSR	22.9±0.3	(22–23)	22.0 ± 0.9	9 (21–23)
AntSR	19.2±1.3	(16–23)	20.3±0.6	(19–21)
PostSR	17.8±0.9	(16–19)	18.0±0.7	(17–19)
PreOc	1.0±	0(1)	1.1±0	3 (1–2)
SupOc	1.0±	0(1)	1.1±0.	3 (1–2)
PostOc	2.0±0.2	2 (2–3)	2.1±0.	3 (2–3)
SubOc	2.2±0.4	4 (2–3)	2.7±0.	5 (2–3)
Ptemp	2±0	(2)	2.0±	0 (2)
Stemp	4.4±0.5	5 (4–5)	4.0±0.	4 (3–5)
SupLab	6.0±	0 (6)	6.0±	0 (6)
InfLab	7.0±	0 (7)	7.1±0.	3 (7–8)
FrL	5.6±0.5 (4.7-	-6.7) N = 20	7.2±0.9	(6.0–9.0)
FrW	3.3±0.3 (2.6-	-4.1) N = 20	3.4±0.5	(2.8–4.4)
FrW/FrL	0.59±0.06 (0.4°	7-0.73) N = 20	0.48 ± 0.02	(0.45–0.52)
SupOcL	5.5±0.5 (4.8-	-6.4) N = 20	6.4±0.7	(5.2–7.4)
SupOcW	3.1±0.3 (2.8-	-3.7) N = 20	3.7±0.5	(3.0–4.6)



FIGURE 9. Type locality habitat of *Acanthophis cryptamydros* **sp. nov.** in Theda Station, Kimberley region, Western Australia (photograph—R.J. Ellis).

Differs from *A. wellsi* by higher midbody scale rows (22–23 vs. 19–21), less prominent dorsal keeling, posterior edge of frontal scale not extending beyond posterior edge of supraoculars (equal to or beyond in *A. wellsi*), and more laterally flared supraocular (absent or less prominent in *A. wellsi*). Differs from melanistic forms of *A. wellsi* by the absence of prominent black coloration on head and black dorsal bands.

Distinguished from *A. antarcticus* by higher average MBSR (23 vs. 21), more ventral scales (125+ vs. 124-), and more prominent anterior dorsal keeling (vs. smooth or very weakly keeled in *A. antarcticus*).

Most similar to *A. rugosus* group in appearance, but differs through higher average ventral scale counts (130 vs. 127), despite considerable overlap in range, lacking dark pigment on the ventrum other than lateral edge (vs. distinct blotching of dark pigment), posterior edge of frontal scale not extending beyond posterior edge of supraoculars (beyond in 13 of 14 specimens, equal in one specimen) and size of lower secondary temporal not larger than sixth supralabial in area (*A. cryptamydros* equal in 13 specimens, smaller in 13 vs. *A. rugosus* equal in 10, larger in 4). See Table 3 for further details.

Comparisons of *A. cryptamydros* **sp nov.** to the *A. rugosus* group are complicated by the likely existence of a number of undescribed species within the latter, which greatly increases variation within this complex. We comment on morphological characters that are useful in distinguishing these two taxa.

Etymology. The specific epithet is modified from the Greek words *kryptos* (cryptic, hidden) and *amydros* (indistinct, dim) in reference to the cryptic nature of the species and its indistinct appearance relative to its surroundings making its presence unknown to predators and prey. Used as a noun in apposition.

Remarks. The *A. rugosus* group is likely to contain a number of undescribed species. Taxonomic resolution of this group will further define differences between individual species within the group and *A. cryptamydros* **sp. nov.** The discovery of *A. cryptamydros* **sp. nov.** as a previously undescribed major lineage within *Acanthophis* highlights the incompleteness of our understanding of phylogenetic structure and species limits within the genus. At the same time, it also highlights the importance of the Kimberley region of Western Australia as a center of endemism (Doughty 2011; Oliver *et al.* 2012; Pepper & Keogh 2014).

Lethal ingestion of the cane toad (*Rhinella marina*) has been documented in previous studies on *Acanthophis* specimens from the Kimberley region and Northern Territory (Phillips & Shine 2007; Hagman *et al.* 2009; Phillips *et al.* 2010; Pearson *et al.* 2014) indicating the species is likely to be at risk of significant decline as cane toads continue to move west across the Kimberley region. A detailed assessment on potential threats to the species including cane toads will identify the need for listing as a species requiring protection under state or federal legislation.

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APPENDIX 1. Specimens used in this study including GenBank accession numbers. Genus abbreviations are as follows: A = Acanthophis, O = Oxyuranus,

Species	Reg. no.	Sex	Locality	State	Lat. (S)	Long. (E)	nd4	cytb	16s	prir	ubn1	Mor.
	QM		Canungra		28°07'45"	153°06'19"						
A. antarcticus	4153 WAM			QLD			KT183462	KT026541	KT026514			
A. antarcticus	2160		Eucla	WA	31°43′	128°54′						•
	WAM											
A. antarcticus	26689 WAM		Araluen	WA	32°08′	116°07′						•
A. antarcticus	26803 WAM		Canning Dam	WA	32°09′	116°08′						•
A. antarcticus	28096		Esperance	WA	33°52′	121°53′						•
0:0:+0x0+0x0	WAIVI		Madura	V/V/	,00°00	176°26'						•
A. antarcticus	37720 WAM		Madura	X X	37 00	120 35						•
A. antarcticus	40197 WAM		Caiguna	WA	32°16′	125°29′						•
A. antarcticus	54251 WAM		Canning Dam Pickering	WA	32°09′	116°07′						•
A. antarcticus	58779 WAM		brook	WA	32°02′	116°06′						•
A. antarcticus	64698 WAM		Canning Dam 10 km SE	WA	32°09′	116°08′						•
(: - ; + · · · · · · · · · · · · · · · · · ·	7000		-	4 / 4 /		700077						

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Re	Reg. no.	Sex	Locality	State	Lat. (S)	Long. (E)	nd4	cytb	16 s	prlr	ubn1	Mor.
WAM	M											
108	108193	ш	Howick Hill 4 km W	WA	33°44′	122°45′						•
W	WAM		Mundaring									
113	113756		Weir	WA	31°58′	116°07′						•
WAM 16569	WAM 165696	ц	Illawarra	WA	32°06'57"	116°08'48"						•
W	WAM											
16569 WAM	165697 WAM	ш	Karragullen	WA	32°07'05″	116°09'08″						•
165	165698	щ	Illawarra	WA	32°09′	116°11'33"						•
WA	WAM											
165	165954	ш	Illawarra	WA	32°10'50"	116°14'35"	KT183456	KT026535	KT026508			•
	WAM		Kunmunya									
A. cryptamydros 5709	6(Mission	WA	15°26′	124°40′						•
WAM	Σ											
A. cryptamydros 106 WA	10628 WAM		Wyndham	WA	15°29′	128°07′						•
A. cryptamydros 112	11241	\supset	Wotjulum	WA	16°11′	123°37′						:
WAIVI A. cryptamydros 13517	WAM 13517A		Yirrkala	W	12°15′	136°53′						•
WAM	Σ											
A. cryptamydros 13517 WAM	13517B WAM		Yirrkala	WA	12°15′	136°53′						•
A. cryptamydros 215	21519		Ranken River	WA	20°04′	137°01′						•
	WAM											
A. cryptamydros 29141 WAM	29141 WAM		Koolan Island	WA	16°09′	123°45′						•
A. cryptamydros 340	34078		Kalumburu	WA	14°18′	126°38′						•

Mor. ... continued on the next page KT026565 KT026562 ubn1 KT026579 KT026581 prlr KT026510 KT026507 **16s** KT026538 KT026534 cytbKT183459 KT183455 nd4 125°44'34" Long. (E) 128°19′ 123°45′ 123°45′ 123°45′ 123°45′ 124°56′ 127°50′ 128°59′ 123°45′ 128°44′ 126°38′ 125°07′ 128°41′ 14°39'46" Lat. (S) 14°18′ 16°09′ 16°09′ 16°09′ 16°09′ 14°36′ 15°20′ 14°00′ 15°54′ 16°09′ 17°14′ 15°47′ 17°51′ State ٨ ٨ ٨ Χ Α ٨ ΑM W۸ ٨ Α× W۸ ٨ ٨ × Surveyors Pool Parry Harbour Bungle Bungle River Nat Res Koolan Island Koolan Island Koolan Island **Koolan Island Koolan Island** Prince Regent Gibson Point, Bigge Island 45 km NNE Locality Kalumburu Halls Creek acksaddle Kununurra Kununurra Nat. Park Sex Σ Σ Σ Σ Σ \supset APPENDIX 1. (continued) Reg. no. 103755 137470 106033 116934 34079 37762 37763 37764 41457 46836 WAM 70690 WAM 70698 81245 WAM WAM WAM 37761 WAM WAM WAM WAM WAM WAM WAM WAM WAM A. cryptamydros Species (P

Mor. • • • • • • KT026566 KT026563 KT026564 KT026561 KT026567 ubn1 KT026580 KT026583 KT026582 KT026578 prlr KT026512 KT026511 KT026513 KT026518 KT026519 KT026517 KT026509 16s KT026544 KT026540 KT026560 KT026546 KT026539 KT026545 KT026537 KT026536 cytb KT183476 KT183460 KT183458 KT183465 KT183477 KT183457 KT183461 nd4 123°45'05" 123°49'49" 124°13'49" 124°13'37" 140°26'50" Long. (E) 126°29'22' 145°29'6" 125°44′ 125°42′ 125°12′ 119°29′ 06°39'11" 16°16'38" 16°08'04" 16°22'21" 14°46'59" 16°22'13" 16°14'17" Lat. (S) 15°05′ 20°17′ 14°50′ 14°49′ State QLD $\frac{1}{2}$ ٨ ٨ ٨ $^{\vee}$ $\stackrel{\mathsf{A}}{\sim}$ $\stackrel{\mathsf{A}}{\sim}$ $^{\vee}$ ٨ $^{\times}$ 9 $\stackrel{\mathsf{A}}{\mathsf{A}}$ F Adelaide River Kuala Kencana **Mertens Falls** Little Mertens Koolan Island NW Molema SE Theda HS Soldsworthy Locality Boongaree 7 km NNW nr Timika) Kimberley unknown Dorat Rd, Wulalam Wulalam Island Island Island Island Asike Cape Falls Sex Σ ≥ ≥ ш ш ш APPENDIX 1. (continued) Reg. no. 145216 172035 168918 141552 171659 172034 174083 164794 165567 WAM WAM WAM WAM WAM WAM WAM WAM 29109 WAM WAM NTM 1774 3286 3214 **≫ ≫ %** A. cryptamydros A. praelongus Species A. pyrrhus A. laevis A. Iaevis (P (P

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Species	Reg. no.	Sex	Locality	State	Lat. (S)	Long. (E)	nd4	cytb	16s	prir	ubn1	Mor.
	WAM											
A. pyrrhus	85117 WAM		Port Hedland	WA	20°19′	118°36′						•
A pyrrhus	91671		Plains HS	δ/M	19°44′	121°24'						•
	WAM		55 km S Anna			- 1 1)
A. pyrrhus	91672		Plains HS	WA	19°44′	121°24′						•
	WAM		30 km SE									
A. pyrrhus	104357		South Hedland	WA	20°25′	118°56′						•
			Kennedy									
	WAM		Range									
A. pyrrhus	123191		National Park	WA	24°30'03"	115°01'03"						•
	WAM		38 km N Port									
A. pyrrhus	124838		Heldand	WA	20°25′	118°50′						•
	WAM		38 km N Port									
A. pyrrhus	124878		Heldand	WA	20°25′	118°50′						•
	WAM											
A. pyrrhus	129434 WAM		Port Hedland	WA	20°19′	118°36′						•
A. pvrrhus	129435		Port Hedland	W	20°19′	118°36′						•
			3 km W))) !						
	WAM		Sandfire									
A. pyrrhus	135628		Roadhouse 143 km S	WA	19°46′	121°03′						•
	WAM		Roebuck Plains									
A. pyrrhus	137991		Roadhouse	WA	19°45′	123°05′						•
	WAM		37 km E South									
A. pyrrhus	141275	ш	Hedland	WA	20°24′	118°56′						•
	WAM		35 km E South									
A. pyrrhus	141276	ш	Hedland	WA	20°24′	118°55′						•

APPENDIX 1. (continued)	(continued	1)										
Species	Reg. no.	Sex	Locality	State	Lat. (S)	Long. (E)	nd4	cytb	16s	prir	ubn1	Mor.
			Carouse Dam,									
	WAM		110 km NE									
A. pyrrhus	146966	щ	Kalgoorlie	WA	30°09'13"	122°21'22"						•
			Carouse Dam,									
	WAM		110 km NE									
A. pyrrhus	154930	Σ	Kalgoorlie	WA	30°12′	122°24′	KT183467	KT026548	KT026521		KT026570	•
	WAM											
A. pyrrhus	156223	Σ	Shay Gap area	WA	20°26'38"	120°00'14"						•
	WAM											
A. pyrrhus	156224	ш	Shay Gap area	WA	20°26'38"	120°00'14"						•
	WAM		Mandora									
A. pyrrhus	162978		Station	WA	19°44′	120°50′						•
			39 km NE									
	WAM		Minilya									
A. pyrrhus	165891	ш	Roadhouse	WA	23°34′	114°17′	KT183466	KT026547	KT026520	KT026585	KT026569	•
			South Alligator									
A. 'rugosus	MTN		River									
group'	9724	щ	Floodplain	N	12°40'59"	132°31′						•
			South Alligator									
A. 'rugosus	MTN	щ	River									
group'	17879		Floodplain	Z	12°40'59"	132°31′						:
			South Alligator									
A. 'rugosus	MTN		River									
group'	17880	щ	Floodplain	N	12°40'59"	132°31′						•
A. 'rugosus	MTN		nr. Nguiu,									
group'	17881	щ	Bathurst Is.	Ν	11°43'54″	130°34'17"						•
			Daly River Rd.,									
A. 'rugosus	MTN		Daly River									
group'	29847	Σ	Region	Z	13°29'34″	131°02'24"	KT183471	KT026551	KT026525	KT026588	KT026573	•
											on out no po	4

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Mor. • • • • • • KT026576 KT183472 KT026552 KT026526 KT026589 KT026574 KT026575 ubn1 KT026553 KT026527 KT026590 KT026591 prlr KT026528 KT026524 KT026522 **16**s KT026554 KT026550 KT026558 cytbKT183470 KT183473 KT183468 nd4 131°15'51" 130°56'60" 131°09'15" 139°30'48" Long. (E) 131°09'34" 131°09'01" 130°05'05" 131°09'37" 135°04′ 13°24'04" 13°29'13" 13°24'06" 13°31'36" 13°23'28" 13°23'37" 20°40'09" Lat. (S) 12°49′ 18°00′ State QLD $\frac{1}{2}$ F \vdash F \vdash F F F 닐 Adelaide River Adelaide River Adelaide River Daly River Rd., Adelaide River Daly River Rd. Road, Darwin Moondarra, Locality Old Bynoe Dorat Rd., Dorat Rd., Dorat Rd., Daly River Daly River Dorat Rd., Anthonys Merauke Region Region Lagoon Region Region Region Region River Sex Σ Σ Σ Σ Σ ≥ Σ APPENDIX 1. (continued) Reg. no. WW 278 29918 34976 35380 NTM 35709 31212 34980 35303 35641 NTM MTM MTM MTM MTM MTM MTM QM 5222 Species A. 'rugosus group' group' group' group' group' group' group' group' group'

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Mor. KT026592 KT026577 KT183454 KT026533 KT026506 KT026584 KT026568 KT026587 KT026572 KT026571 ubn1 KT026586 prlr KT026516 KT026530 KT026529 KT026515 KT026523 **16**s KT026556 KT026549 KT026542 KT026543 KT026555 cytbKT183469 KT183463 KT183464 KT183475 KT183474 nd4 115°31'20" 115°31'31" 120°27'20" Long. (E) 138°5'0" 114°10′ 116°19′ 114°07′ 113°49′ 114°02′ 118°48′ 119°44′ 116°19′ 22°23'47" 22°07'11" 21°13'04" 19°16'0" Lat. (S) 21°48′ 21°56′ 22°20′ 22°03′ 21°39′ 21°39′ 22°03′ 23°21′ State QLD ٨ ٨ ٨ ٨ $^{\mathsf{W}}$ ٨ ٨ ٨ ٨ ٨ Α× F Adelaide River Newman area **Mount Minnie** Vlaming Head Pannawonica Pannawonica **Yardie Creek** Meentheena flood plains Locality -ighthouse Roadhouse Roadhouse **Fablelands** Nanutarra Merauke Exmouth Shothole Munjina Canyon Mouth Barkly area area Sex ≥ Σ APPENDIX 1. (continued) Reg. no. QM 516 113377 113378 119367 129559 139366 136097 139188 93215 26759 WAM WAM 13437 19674 WAM 61495 WAM WAM WAM WAM WAM WAM 3295 SAM Species A. 'rugosus A. 'rugosus A. 'rugosus A. wellsi group' group' group'

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APPENDIX 1. (continued)	[continued	(1										
Species	Reg. no.	Sex	Locality	State	Lat. (S)	Long. (E)	nd4	cytb	16s	prir	ubn1	Mor.
			2km NE									
A wellsi	WAINI 142608	Σ	Snotnole Canvon	ΜA	22°02′	114°02′						•
		:	19 km N])]	 - -						1
	WAM		Nanutarra									
A. wellsi	146967	ш	Roadhouse	WA	22°22′	115°30′						•
	WAM		3 km NE									
A. wellsi	151150	ш	Mound Minnie	٨٨	22°05′	115°34′						•
	WAM		100 km NW									
A. wellsi	151282		Wittenoom	٨	21°44'34"	117°19'09"						•
	WAM		Shothole									
A. wellsi	154974	щ	Canyon	۸۸	22°03'13"	114°01'11"						•
	WAM		Chichester									
A. wellsi	156324	Σ	Range	WA	22°13'49"	118°58'53"						•
	WAM		Chichester									
A. wellsi	156325	Σ	Range	WA	22°13'46"	119°00'05"						•
	WAM											
A. wellsi	157563	Σ	Robe River	WA	21°07'57"	115°52'10"						•
	WAM											
A. wellsi	163236		Jinayri Mine	٨٨	23°01'01"	119°10'23"						•
	WAM		Mt Stewart-									
A. wellsi	165176		Wyloo area	۸								•
	WAM		7 km SE									
A. wellsi	170589		Peedamulla	٨٨	21°52'15"	115°40'47"						•
	WAM		Yandagee									
A. wellsi	170708		Gorge	WA	21°38'36"	116°05'48"						•
O. scutellatus	WW 274		Merauke	IND			AY340787	KT026557	KT026532			
			Bamustu,									
			Western									
P. papuanus	WW 844		Province	PNG			AY340144	KT026559	KT026531			